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L5 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN
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RN 26194-60-5 REGISTRY

ED Entered STN: 16 Nov 1984

CN 2,3-Naphthalenedimethanol, 1α -(3,4-dihydroxyphenyl)- 1,2 α ,3 β ,4-tetrahydro-6,7-dimethoxy-, tetraacetate (8CI) (CA INDEX NAME)

OTHER NAMES:

CN Isotaxiresinol 6-methyl ether tetraacetate

MF C28 H32 O10

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

Currently available stereo shown.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L5 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN
- RN 26194-57-0 REGISTRY
- ED Entered STN: 16 Nov 1984
- CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

OTHER CA INDEX NAMES:

- CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, $(1\alpha,2\beta,3\alpha)$ -
- CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, stereoisomer (8CI)

OTHER NAMES:

- CN (+)-Isotaxiresinol
- CN Isotaxiresinol
- FS STEREOSEARCH
- MF C19 H22 O6
- LC STN Files: AGRICOLA, BEILSTEIN*, BIOSIS, CA, CAPLUS, CHEMCATS, IPA, NAPRALERT, TOXCENTER, USPATFULL

(*File contains numerically searchable property data)

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

35 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

35 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 3 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN 23141-17-5 REGISTRY L5

RN

ED Entered STN: 16 Nov 1984

2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-6,7dimethoxy-, $(1\alpha, 2\beta, 3\alpha)$ - (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

2,3-Naphthalenedimethanol, 1α -(3,4-dihydroxyphenyl)-

1, 2α , 3β , 4-tetrahydro-6, 7-dimethoxy- (8CI)

OTHER NAMES:

CN Isotaxiresinol 6-methyl ether

FS STEREOSEARCH

C20 H24 O6 ${
m MF}$

LC STN Files: BEILSTEIN*, BIOSIS, CA, CAPLUS (*File contains numerically searchable property data)

Relative stereochemistry. Currently available stereo shown.

ОН ОН OMe HO R R HO OMe

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
=> s demethylisolariciresinol
L6
             2 DEMETHYLISOLARICIRESINOL
=> d 1-2
     ANSWER 1 OF 2 REGISTRY COPYRIGHT 2009 ACS on STN
L6
RN
     349150-66-9 REGISTRY
ΕD
     Entered STN: 27 Jul 2001
     2,3-Naphthalenedimethanol, 1,2,3,4-tetrahydro-6,7-dihydroxy-1-(4-hydroxy-3-
CN
     methoxyphenyl)-, (1S, 2R, 3R)- (CA INDEX NAME)
OTHER NAMES:
CN
     (-)-3-Demethylisolariciresinol
FS
     STEREOSEARCH
     C19 H22 O6
MF
SR
     CA
LC
     STN Files:
                  CA, CAPLUS, TOXCENTER
Absolute stereochemistry. Rotation (-).
          ОН
                 OMe
                       ОН
НО
        R
        R
HO
                       ОН
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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6 REFERENCES IN FILE CA (1907 TO DATE) 6 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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ANSWER 2 OF 2 REGISTRY COPYRIGHT 2009 ACS on STN
L6
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RN 349150-65-8 REGISTRY

ΕD Entered STN: 27 Jul 2001

1,2-Benzenediol, 4-[(1S,2R,3R)-1,2,3,4-tetrahydro-7-hydroxy-3-CN (hydroxymethyl)-2-[(1-hydroxy-1-methylethoxy)methyl]-6-methoxy-1naphthalenyl] - (CA INDEX NAME)

OTHER NAMES:

CN (-)-3'-Demethylisolariciresinol-9'-hydroxyisopropyl ether

FS STEREOSEARCH

C22 H28 O7 MF

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 5 REFERENCES IN FILE CA (1907 TO DATE)
- 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=>

Uploading C:\Documents and Settings\byongkwon\My Documents\osteo-1.str

L7 STRUCTURE UPLOADED

=> s sss samp 17

SAMPLE SEARCH INITIATED 09:36:22 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 230 TO ITERATE

100.0% PROCESSED 230 ITERATIONS 1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 3691 TO 5509 PROJECTED ANSWERS: 1 TO 80

L8 1 SEA SSS SAM L7

=> d

L8 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2009 ACS on STN

RN 19856-53-2 REGISTRY

ED Entered STN: 16 Nov 1984

CN 2-Naphthalenecarboxylic acid, $1-(3,4-dihydroxy-5-methoxyphenyl)-1,2,3,4-tetrahydro-2,3,7-trihydroxy-3-(hydroxymethyl)-6-methoxy-, monopotassium salt, <math>[1S-(1\alpha,2\alpha,3\beta)]-(9CI)$ (CA INDEX NAME)

OTHER CA INDEX NAMES:

- CN 2-Naphthoic acid, 1-(3,4-dihydroxy-5-methoxyphenyl)-1,2,3,4-tetrahydro-2,3,7-trihydroxy-3-(hydroxymethyl)-6-methoxy-, monopotassium salt, (1S,2S,3R)- (8CI)
- CN Plicatic acid, potassium salt (6CI)
- FS STEREOSEARCH
- DR 23486-98-8
- MF C20 H22 O10 . K
- LC STN Files: BEILSTEIN*, CA, CAPLUS, TOXCENTER

(*File contains numerically searchable property data)

CRN (16462-65-0)

Absolute stereochemistry.

K

- 3 REFERENCES IN FILE CA (1907 TO DATE)
- 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s sss 17 full

FULL SEARCH INITIATED 09:36:51 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED -4362 TO ITERATE

100.0% PROCESSED 13 ANSWERS 4362 ITERATIONS

SEARCH TIME: 00.00.01

L9 13 SEA SSS FUL L7

=> file caplus

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 584.96 585.18

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FILE COVERS 1907 - 22 May 2009 VOL 150 ISS 22 (20090521/ED) FILE LAST UPDATED: 21 May 2009 REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

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CAplus now includes complete International Patent Classification (IPC)
reclassification data for the third quarter of 2008.
CAS Information Use Policies apply and are available at:
http://www.cas.org/legal/infopolicy.html
This file contains CAS Registry Numbers for easy and accurate
=> s 19
L10
            82 L9
=> s 110 and (osteo? or osteoporosis OR "Bone resorption" OR "Bone resorption" OR
"Bone, disease" or bone or osteoclast)
         98515 OSTEO?
         25271 OSTEOPOROSIS
        249972 "BONE"
         25462 "BONES"
        257144 "BONE"
                 ("BONE" OR "BONES")
         36528 "RESORPTION"
          1053 "RESORPTIONS"
         37399 "RESORPTION"
                 ("RESORPTION" OR "RESORPTIONS")
         15552 "BONE RESORPTION"
                 ("BONE"(W) "RESORPTION")
        249972 "BONE"
         25462 "BONES"
        257144 "BONE"
                 ("BONE" OR "BONES")
         36528 "RESORPTION"
          1053 "RESORPTIONS"
         37399 "RESORPTION"
                 ("RESORPTION" OR "RESORPTIONS")
         15552 "BONE RESORPTION"
                 ("BONE"(W) "RESORPTION")
        249972 "BONE"
         25462 "BONES"
        257144 "BONE"
                 ("BONE" OR "BONES")
       1152627 "DISEASE"
        317869 "DISEASES"
       1293574 "DISEASE"
                 ("DISEASE" OR "DISEASES")
         23700 "BONE, DISEASE"
                 ("BONE"(W)"DISEASE")
        249972 BONE
         25462 BONES
        257144 BONE
                 (BONE OR BONES)
         10702 OSTEOCLAST
          7089 OSTEOCLASTS
         12451 OSTEOCLAST
                 (OSTEOCLAST OR OSTEOCLASTS)
L11
             2 L10 AND (OSTEO? OR OSTEOPOROSIS OR "BONE RESORPTION" OR "BONE
               RESORPTION" OR "BONE, DISEASE" OR BONE OR OSTEOCLAST)
=> s 110 and (isotaxiresional or taxus or yunnanesis or taxus chinenesis or chinese
yew or isolariciresinol or taxus baccata or english yew or yew)
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0 ISOTAXIRESIONOL

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19 YUNNANESIS
          3014 TAXUS
            14 CHINENESIS
             1 TAXUS CHINENESIS
                 (TAXUS(W)CHINENESIS)
        119204 CHINESE
            32 CHINESES
        119227 CHINESE
                 (CHINESE OR CHINESES)
          1854 YEW
            42 YEWS
          1864 YEW
                (YEW OR YEWS)
            51 CHINESE YEW
                (CHINESE(W)YEW)
           250 ISOLARICIRESINOL
          3014 TAXUS
           827 BACCATA
           590 TAXUS BACCATA
                (TAXUS(W)BACCATA)
         12830 ENGLISH
          1854 YEW
            42 YEWS
          1864 YEW
                (YEW OR YEWS)
            13 ENGLISH YEW
                (ENGLISH(W)YEW)
          1854 YEW
            42 YEWS
          1864 YEW
                 (YEW OR YEWS)
L12
            32 L10 AND (ISOTAXIRESIONOL OR TAXUS OR YUNNANESIS OR TAXUS CHINENE
               SIS OR CHINESE YEW OR ISOLARICIRESINOL OR TAXUS BACCATA OR ENGLI
               SH YEW OR YEW)
=> focus
PROCESSING COMPLETED FOR L12
             32 FOCUS L12 1-
=> d ibib abs hitstr 1-32
L13 ANSWER 1 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER:
                       1952:54579 CAPLUS
DOCUMENT NUMBER:
                        46:54579
ORIGINAL REFERENCE NO.: 46:9086b-i,9087a-b
                         Isotaxiresinol (3'-dimethylisolariciresinol), a new
TITLE:
                         lignan extracted from the heartwood of the
                         English yew, Taxus
                         baccata
                         King, F. E.; Jurd, L.; King, T. J.
AUTHOR(S):
CORPORATE SOURCE:
                         Univ. Nottingham, UK
SOURCE:
                         Journal of the Chemical Society (1952) 17-24
                         CODEN: JCSOA9; ISSN: 0368-1769
DOCUMENT TYPE:
                         Journal
                         Unavailable
LANGUAGE:
OTHER SOURCE(S):
                        CASREACT 46:54579
GΙ
    For diagram(s), see printed CA Issue.
     The heartwood (I) of the English yew (Taxus
    baccata) (1000 g.) was boiled 3 times (each 2 h.) with H2O, the
     extract concentrated, the filtrate from the dark brown resinous precipitate
(II) extracted 7
```

3014 TAXUS

times with ether, and the residue from the ether boiled with AcOEt, giving 5.4 g. isotaxiresinol (III), m. 171°; II becomes partly crystalline when treated with 40% AcOH and, crystallized from 2 N aqueous AcOH, gives an addnl. 3 g. of III (total yield, 1% of undried I). I (200 g.), extracted 2 days with ether, and the oily residue from the extract refluxed with petr. ether and crystallized from AcOEt, gives 0.4 g. III; no evidence of an isomer of III was obtained. With Me2SO4 in 2 N NaOH at 60° (final heating for 10 min. on a steam bath), III yields the tri-Me ether (IV) (isolariciresinol dl-Me ether), m. 167-8°, $[\alpha]$ D18 19° (CHCl3) (Haworth and Kelly, C.A. 31, 3930.7). III (1 q.), 5 cc. EtI, and 20 cc. Me2CO containing 2 g. anhydrous K2CO3, refluxed 30 h., give 0.75 g. of the tri-Et ether (V), m. 140° . IV (0.2 g.) and 0.8 g. KHSO4, heated 0.5 h. at $180-90^{\circ}$, give anhydroisotaxiresinol tri-Me ether (anhydroisolariciresinol di-Me ether), m. 149.5° the corresponding tri-Et ether m. 132.5-3° (diacetate, m. 89.5°; dibenzoate, m. 125°). Oxidation of V with HNO3 gives 4,5,1,2-(O2N)2C6H2(OEt)2; 0.2 g. IV yields 0.04 g. 4,5,1,2-(O2N) 2C6H2 (OMe) 2. V (0.35 g.) in 40 cc. boiling Me2CO, treated (2 h.) with 1.5 g. powdered KMnO4, gives 2-(3,4-diethoxybenzoyl)-4-ethoxy-5-methoxybenzoic acid (VI), m. 173° (Me ester, m. 111°); VI results also on refluxing V with K2Cr207 in AcOH (3 h.). The structure of VI was established by the following synthesis. 3,4-(HO)2C6H3CHO (13.8 g.), slowly treated (10 min.) with 38.5 g. Et2SO4 and 18.4 g. KOH in 50 cc. H2O, gives 79% $3,4-(EtO)\,2C6H3CHO$ (VII), b. $280-2^{\circ}$ (semicarbazone, pale yellow, m. 175°). VII (33 g.) with 43 g. KMnO4 in 800 cc. H2O (boiled 5 min.) gives 88% 3,4-(EtO)2C6H3CO2H, m. 166-7° (amide, m. 183.5°). 4,3-EtO(MeO)C6H3Me (5.45 g.) and 7.5 g. 3,4-(EtO)2C5HCOCl (VIII) in 30 g. PhNO2 (ice bath), treated (10 min.) with 11.1 g. AlC13 and kept overnight, give 40% 3',4',5-triethoxy-4-methoxy-3-methylbenzophenone, m. 115°; this is not oxidized by KMnO4; CrO3 in AcOH (with or without H2SO4) gives 3,6,7-triethoxy-2-methoxyanthraquinone (IX), bright yellow, m. 225°, and a small quantity of an unidentified acid; K2Cr2O7 in boiling 75% AcOH gives a small quantity of IX, an alkali-insol. compd, m. 152°, and a small yield of an acid, probably VI, but difficult to purify. 4,3-EtO(MeO)C4H3CHO (25 g.) and 35 g. CH2(CO2H)2 in 50 cc. C5H5N and 2.5 cc. piperidine, heated 1 h. on a steam bath and refluxed 15 min., the acid in N NaOH reduced with 700 g. 3% Na-Hq, and the product (28 g.) esterified with MeOH-HCl (refluxed 15 h.), give 27 g. Me β -(4-ethoxy-3-methoxyphenyl)propionate (X), m. 37-8°. X and N2H4.H2O, heated 3 h. on the steam bath, give the hydrazide m. 123°; this yields 4,3-EtO(MeO)C6H3CH2NH2 (XI). XI (4 g.) in 10 cc. C5H5N, gradually treated with 5.2 g. VIII in 25 cc. C6H6 and heated 15 min. on the steam bath, gives 86% 3,4-diethoxy-N-(4-ethoxy-3methoxyphenethyl)benzamide (XII), m. 148.5°. XII (7.5 g.), 15 g. POC13, and 30 cc. PhMe, refluxed 2 h., give the HCl salt, yellow, m. 123° , of 1-(3,4-diethoxyphenyl)-7-ethoxy-3,4-dihydro-6methoxyisoquinoline, pale brown, m. 115-15.5°, 82%; methiodide (XIII), with 0.5 mol. H2O, yellow, m. 192°. XIII (2 g.) in 12 cc. MeI, shaken 1.5 h. with 80 cc. N NaOH at room temperature, gives a nearly quant. yield of [2-(3,4-diethoxybenzoyl)-4-ethoxy-5methoxyphenethyl]trimethylammonium iodide (XIV), m. 159° (picrate, yellow, m. 171.5°). XIV (1.8 g.) in 50 cc. 2 N NaOH, heated 15 min. on a steam bath, gives 56% 3β , 4β , 5-triethoxy-4-methoxy-2vinylbenzophenone (XV), m. $118-18.5^{\circ}$. Oxidation of XV with KMnO4 in Me2CO gives VI; VI with concentrated H2SO4 yields IX. I (1200 g.), extracted twice with cold petr. ether and twice with ether, and the dried residue extracted

twice with cold H2O, gives 2.5 g. III; the filtrate from the III yields

0.51 g. (0.04%) of sequoyitol, m. 237° (Sherrard and Kurth, C.A. 23, 5469). Color reactions indicate the absence of flavones, flavanones, and flavonolones in I.

IT 477-72-5P, 2,3-Naphthalenedimethanol,

1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-

26194-57-0P, Isotaxiresinol

RL: PREP (Preparation)
(preparation of)

RN 477-72-5 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy- (CA INDEX NAME)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 2 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1973:111031 CAPLUS

DOCUMENT NUMBER: 78:111031

ORIGINAL REFERENCE NO.: 78:17819a,17822a

TITLE: Taxiresinol, a new lignan in the heartwood of

Taxus baccata

AUTHOR(S): Mujumdar, R. B.; Srinivasan, R.; Venkataraman, K.

CORPORATE SOURCE: Natl. Chem. Lab., Poona, India

SOURCE: Indian Journal of Chemistry (1972), 10(7), 677-80

CODEN: IJOCAP; ISSN: 0019-5103

DOCUMENT TYPE: Journal LANGUAGE: English

GI For diagram(s), see printed CA Issue.

AB A new Lignan, taxiresinol (I) was isolated from Indian **Taxus**baccata, in addition to the known isotaxiresinol (II) and
3,4-MeO(HO)C6H3-CH2CH(CH2OH)CH(CH2OH)CH2-C6H3(OH)OMe-4,3
(secoisolariciresinol) (III). Crystalline acetonides were prepared from the latter two lignans by treatment of acetone solns. with anhydrous copper sulfate, but III gave 3,4-divanillytetrahydrofuran (IV), isolated earlier from Picea excelsa, when copper sulfate is replaced by perchloric acid. The structure (I) is supported by NMR and mass spectral data.

IT **26194-57-0**

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(of Taxus baccata)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 3 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1994:529988 CAPLUS

DOCUMENT NUMBER: 121:129988

ORIGINAL REFERENCE NO.: 121:23393a,23396a

TITLE: A lignan from needles of Himalayan Taxus

baccata

AUTHOR(S): Das, B.; Takhi, M.; Srinivas, K. V. N. S.; Yadav, J.

S.

CORPORATE SOURCE: Org. Chem. Div.-I, Indian Inst. Chem. Technol.,

Hyderabad, 500 007, India

SOURCE: Phytochemistry (1994), 36(4), 1031-3

CODEN: PYTCAS; ISSN: 0031-9422

DOCUMENT TYPE: Journal LANGUAGE: English

AB A new lignan, 4-0-methyl-3'-0-demethyl-(-)-secoisolariciresinol, was

isolated from the needles of Himalayan yew, Taxus

baccata. The structure of the compound was established from its spectral data and chemical reactions.

IT 26194-57-0, Isotaxiresinol
 RL: BIOL (Biological study)

(from Taxus baccata)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

L13 ANSWER 4 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2000:780181 CAPLUS

DOCUMENT NUMBER: 135:270031

TITLE: Studies on the Himalayan yew Taxus

wallichiana: part VII - the taxoids and phenolic

constituents of the roots of Taxus

wallichiana. [Erratum to document cited in

CA131:308822]

AUTHOR(S): Chattopadhyay, S. K.; Kulshrestha, M.; Tripathi, V.;

Saha, G. C.; Sharma, R. P.; Mehta, V. K.

CORPORATE SOURCE: Central Institute of Medicinal and Aromatic Plants,

Lucknow, 226 015, India

SOURCE: Indian Journal of Chemistry, Section B: Organic

Chemistry Including Medicinal Chemistry (2000),

39B(7), 562

CODEN: IJSBDB; ISSN: 0376-4699

PUBLISHER: National Institute of Science Communication, CSIR

DOCUMENT TYPE: Journal LANGUAGE: English

AB The structure published on page 701 is in error; the correct structure is given.

IT **26194-57-0**, Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(from roots of **Taxus** wallichiana (Erratum))

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

L13 ANSWER 5 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1999:580819 CAPLUS

DOCUMENT NUMBER: 131:308822

TITLE: Studies on the Himalayan yew Taxus

wallichiana: part VII - the taxoids and phenolic

constituents of the roots of Taxus

wallichiana

AUTHOR(S): Chattopadhyay, S. K.; Kulshrestha, M.; Tripathi, V.;

Saha, G. C.; Sharma, R. P.; Mehta, V. K.

CORPORATE SOURCE: Central Institute of Medicinal and Aromatic Plants,

Lucknow, 226 015, India

SOURCE: Indian Journal of Chemistry, Section B: Organic

Chemistry Including Medicinal Chemistry (1999),

38B(6), 701-704

CODEN: IJSBDB; ISSN: 0376-4699

PUBLISHER: National Institute of Science Communication, CSIR

DOCUMENT TYPE: Journal LANGUAGE: English

AB The systematic investigation on the roots of **Taxus** wallichiana has resulted in the isolation of nine taxoids – taxol, baccatin III, baccatin IV, taxusin, a C-14 oxygenated taxoid, $5,1\beta$ -hydroxybaccatin I, pentaacetoxy taxadiene, a dibenzoylated rearranged taxoid, 7-xylosyl-10-deacetyl-taxol C and three phenolic compds. (-)-seco-isolariciresinol, taxiresinol and isotaxiresinol. The compds. have been characterized on the basis of their spectral characteristics. The occurrence of taxoid 9 in the roots of the plant is quite significant. The distribution of the above compds. in other parts of the plant are also summarized.

IT **26194-57-0**, Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(from roots of Taxus wallichiana)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 6 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1994:27463 CAPLUS

DOCUMENT NUMBER: 120:27463
ORIGINAL REFERENCE NO.: 120:5093a,5096a

TITLE: Phenolics from needles of Himalayan Taxus

baccata

AUTHOR(S): Das, B.; Takhi, M.; Srinivas, K. V. N. S.; Yadav, J.

S.

CORPORATE SOURCE: Org. Chem. Div. I, Indian Inst. Chem. Technol.,

Hyderabad, 500 007, India

SOURCE: Phytochemistry (1993), 33(6), 1489-91

CODEN: PYTCAS; ISSN: 0031-9422

DOCUMENT TYPE: Journal LANGUAGE: English

GΙ

AB Chemical investigation on the needles of the Himalayan **yew** resulted in the isolation of several phenolic compds., including 3-demethyl-(-)-secoisolariciresinol (I), a new lignan, and taxuside (II), a new phenolic glucoside. Th structures of the new compds. were derived from their spectral data and chemical transformations.

IT **26194-57-0**, Isotaxiresinol

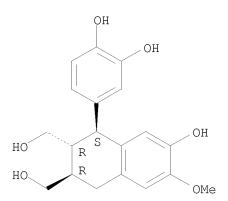
RL: BIOL (Biological study)

(from Himalayan yew needles)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



L13 ANSWER 7 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2001:398696 CAPLUS

DOCUMENT NUMBER: 135:149914

TITLE: Important phenolic constituents of the Himalayan

Yew Taxus wallichiana

AUTHOR(S): Chattopadhyay, S. K.; Kulshrestha, M.; Tripathi, V.;

Sashidhara, K. V.; Kumar, Sushil

CORPORATE SOURCE: Central Institute of Medicinal and Aromatic Plants,

Lucknow, 226 015, India

SOURCE: Journal of Medicinal and Aromatic Plant Sciences

(2000), 22(1B), 710-714

CODEN: JMASF6

PUBLISHER: Central Institute of Medicinal and Aromatic Plants

DOCUMENT TYPE: Journal LANGUAGE: English

AB A systematic chemical investigation of the leaves, stem bark, heartwood and

roots of Taxus wallichiana has resulted in the isolation of

several phenolic compds. i.e. (-) betuligenol, betuloside, (+) catechin,

(-) secoisolariciresinol, taxiresinol and isotaxiresinol.

IT 26194-57-0P, Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation)

(important phenolic constituents of Himalayan Yew

Taxus wallichiana)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 8 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1997:485598 CAPLUS

DOCUMENT NUMBER: 127:173864

ORIGINAL REFERENCE NO.: 127:33613a,33616a

TITLE: The taxoids and the phenolic constituents of the

heartwood of the Himalayan yew Taxus

wallichiana

AUTHOR(S): Chattopadhyay, S. K.; Kulshrestha, M.; Saha, G. C.;

Sharma, R. P.; Jain, S. P.; Kumar, Sushil

CORPORATE SOURCE: Central Institute of Medicinal and Aromatic Plants,

Lucknow, 226015, India

SOURCE: Journal of Medicinal and Aromatic Plant Sciences

(1997), 19(1), 17-21

CODEN: JMASF6

PUBLISHER: Central Institute of Medicinal and Aromatic Plants

DOCUMENT TYPE: Journal LANGUAGE: English

AB Four taxoids were isolated from the heartwood of T. wallichiana: taxusin, a C-14 oxygenated taxoid, a dibenzoylated rearranged taxoid, and a rare taxol xyloside derivative Also isolated were 3 lignans: taxiresinol, isotaxiresinol, and (-)-secoisolariciresinol. The absolute stereochem. of (-)-secoisolariciresinol was established by x-ray crystallog.

IT **26194-57-0**, Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(taxoids and phenolic constituents of heartwood of **Taxus** wallichiana)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 9 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:66156 CAPLUS

DOCUMENT NUMBER: 143:149902

TITLE: Constituents from the roots of Taxus

cuspidata

AUTHOR(S):

CORPORATE SOURCE:

Kawamura, Fumio; Ohira, Tatsuro; Kikuchi, Yoshinari

Department of Forest Chemistry, Forestry and Forest

Products, Research Institute, Tsukuba, 305-8687, Japan

SOURCE: Journal of Wood Science (2004), 50(6), 548-551

CODEN: JWSCFG; ISSN: 1435-0211

PUBLISHER: Springer Tokyo

DOCUMENT TYPE: Journal LANGUAGE: English

The known propelargonidin, afzelechin- $(4\alpha\rightarrow 8)$ -afzelechin (1), the known lignans 7'-hydroxynortrachelogenin (2), epinortrachelogenin (3), nortrachelogenin (4), hydroxymatairesinol (5), allohydroxymatairesinol (6), matairesinol (7), oxomatairesinol (8), and isotaxiresinol (9), and the known taxoids taxinine M (10), taxayuntin (11), and 10-deacetyltaxol (12), and 10-deacetylbaccatin III (13) were isolated from the roots of **Taxus** cuspidata (Japanese **yew**, Taxaceae). The propelargonidin was isolated from **Taxus** spp. for the first time, and was detected in the roots, bark, and twigs.

IT **26194-57-0P**, Isotaxiresinol

RL: BSU (Biological study, unclassified); NPO (Natural product occurrence); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation)

(isolation and characterization of constituents from the roots of

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Taxus cuspidata)
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RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 10 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1985:565998 CAPLUS

DOCUMENT NUMBER: 103:165998

ORIGINAL REFERENCE NO.: 103:26567a,26570a

TITLE: Constituents of the heartwood of Taiwan yew

AUTHOR(S): Liu, Ching Long; Lin, Yuan Chuan; Lin, Yuh Mei; Chen,

Fa Ching

CORPORATE SOURCE: Dep. Chem., Natl. Taiwan Univ., Taipei, Taiwan

SOURCE: Taiwan Kexue (1984), 38(3), 119-25

CODEN: TKHSAU; ISSN: 0371-845X

DOCUMENT TYPE: Journal LANGUAGE: Chinese

AB The heartwood of Taiwan yew (Taxus mairei) is a folk

drug for treating diabetes. From the hexane-soluble part of the MeOH extract

of

this heartwood, alkanes, fatty acids (C24-26), β -sitosterol

[83-46-5], taxinine [3835-52-7], and taxusin [19605-80-2] were isolated.

From the CHCl3-soluble part of the MeOH extract, vanillin [121-33-5],

coniferaldehyde [458-36-6], α -conidendrin [518-55-8],

(-)-secoisolariciresinol [29388-59-8], meso-secoisolariciresinol

[57759-55-4], taxa-4(20),11-diene-5 α ,9 α ,10 β ,13 α -

tetrol- 9α , 10β -diacetate [27854-00-8], isotaxiresinol

26194-57-0], and a new biphenyl were isolated by silica gel column chromatog. eluting with EtOAc/hexane.

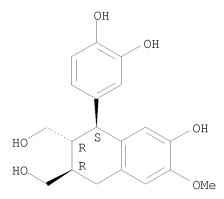
IT **26194-57-0**

RL: BIOL (Biological study)

(of Taxus mairei heartwood)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)



L13 ANSWER 11 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2007:1012507 CAPLUS

DOCUMENT NUMBER: 147:465158

TITLE: Antioxidant activity of polyphenols from the far-east

plant Taxus cuspidata

AUTHOR(S): Veselova, M. V.; Fedoreev, S. A.; Vasilevskaya, N. A.;

Denisenko, V. A.; Gerasimenko, A. V.

CORPORATE SOURCE: Pacific Institute of Bioorganic Chemistry, Far-East

Division, Russian Academy of Sciences, Vladivostok,

Russia

SOURCE: Pharmaceutical Chemistry Journal (2007), 41(2), 88-93

CODEN: PCJOAU; ISSN: 0091-150X

PUBLISHER: Springer
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Phenolic components of the wood and roots of far-east yew (
Taxus cuspidata) have been isolated and investigated. Four lignans [(+)-taxiresinol, (+)-isotaxiresinol, (+)-isolariciresinol , (-)-secoisolariciresinol] and two catechins [(+)-catechin, (-)-epicatechin] were identified using spectroscopic techniques. HPLC data showed that wood, bark, roots, stems, and needles of the plant contain different amts. of lignans and catechins. The antioxidant and radical-scavenging activities of the polyphenols were evaluated on two in vitro model systems.

IT **26194-57-0P**, (+)-Isotaxiresinol

RL: BSU (Biological study, unclassified); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); PREP (Preparation) (antioxidant activity of polyphenols from the far-east plant

Taxus cuspidata)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

CAPLUS COPYRIGHT 2009 ACS on STN L13 ANSWER 12 OF 32

ACCESSION NUMBER: 1992:148178 CAPLUS

DOCUMENT NUMBER: 116:148178

ORIGINAL REFERENCE NO.: 116:24961a,24964a

TITLE: Constituents of the heartwood of Taiwan Yew.

Part IV. Isolation of 1,4-p-methanediol and

1-dehydroxybaccatin-IV

Chuang, L. C.; Chen, K. J.; Lin, Y. S.; Chen, F. C. AUTHOR(S):

CORPORATE SOURCE: Dep. Chem., Tamkang Univ., Tamsui, Taiwan

Huaxue (1990), 48(4), 275-80 SOURCE:

CODEN: HUHSA2; ISSN: 0441-3768

DOCUMENT TYPE: Journal LANGUAGE: Chinese

AΒ The heartwood of Taiwan Yew (Taxus maitei) contained

long chain alkanes, long chain esters, β -sitosterol, taxusin,

taxa-4(20), 11-diene- 5α , 9α , 10β , 13α -tetrol- 9α , 10β -diacetate, taxa-4(20), 11-diene-

 2α , 5α , 7β , 10β -tetrol- 5α , 7β , 10β -

triacetate- 2α -methylbutyrate, secoisolariciresinol,

isotaxiresinol, 1-dehydroxybaccatin IV, (+)-dihydroquercetin, D-sesamin,

4-hydroxysesamin, cis-terpin and 1,4-p-methanediol.

ΙT **26194-57-0**, Isotaxiresinol

RL: BIOL (Biological study)

(from Taxus mairei heartwood)

RN 26194-57-0 CAPLUS

2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-1,2,3,4-tetrahydro-CN hydroxy-6-methoxy-, (1S, 2R, 3R)- (CA INDEX NAME)

L13 ANSWER 13 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1982:578717 CAPLUS

DOCUMENT NUMBER: 97:178717

ORIGINAL REFERENCE NO.: 97:29835a,29838a

TITLE: Studies on the lignans of Zi Shan (Taxus

cuspidata Sieb et Zucc.)

AUTHOR(S): Zhou, Youzuo; Yu, Chaomei; Zhu, Yuanlong

CORPORATE SOURCE: Inst. Pharmacol., Zhejiang Acad. Public Hyg., Peop.

Rep. China

SOURCE: Zhongcaoyao (1982), 13(4), 1-2

CODEN: CTYAD8; ISSN: 0253-2670

DOCUMENT TYPE: Journal LANGUAGE: Chinese

AB Three crystalline components, A, B, and D, were isolated and crystallized from

an

EtOH extract of powdered T. cuspidata wood. Components A and B were yellow needle crystals, whereas D was a white granular crystal. Results from UV, IR, NMR, and mass spectral anal., as well as derivative syntheses revealed components B and D to be isotaxiresinol and isolariciresinol, resp. With the exception of the m.p. difference between components A and B recrystd. from aqueous MeOH, all the physicochem. and other characteristics of A were identical to those of B, indicating that A is an allomorphic isomer of B. None of these components showed antitumor activity, indicating that A, B, and D are not the antitumor substances in T. cuspidata.

IT 26194-57-0

RL: BIOL (Biological study)

(from **Taxus** cuspidata)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 14 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1969:410304 CAPLUS

DOCUMENT NUMBER: 71:10304
ORIGINAL REFERENCE NO.: 71:1883a,1886a

TITLE: Taxus heartwood constituents
AUTHOR(S): Erdtman, Holger; Tsuno, K.

CORPORATE SOURCE: Roy. Inst. Technol., Stockholm, Swed.

SOURCE: Phytochemistry (Elsevier) (1969), 8(5), 931-2

CODEN: PYTCAS; ISSN: 0031-9422

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The heartwood of several **Taxus** species was investigated. All contain a series of 6 lignans also occurring in Fitzroya cupressoides. The neutral constituents, however, are different from those of Fitzroya.

IT **26194-57-0**

RL: BIOL (Biological study)

(of Taxus, taxonomy in relation to)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 15 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:205569 CAPLUS

DOCUMENT NUMBER: 146:12699

TITLE: Chemical constituents of cultured Taxus

mairei (I)

AUTHOR(S): Chen, Xue-Ying; Liang, Jing-Yu

CORPORATE SOURCE: Department of Phytochemistry, China Pharmaceutical

University, Nanjing, 210009, Peop. Rep. China

SOURCE: Zhongguo Tianran Yaowu (2006), 4(1), 52-57

CODEN: ZTYHA7; ISSN: 1672-3651 PUBLISHER: Zhongguo Tianran Yaowu Bianjibu

DOCUMENT TYPE: Journal LANGUAGE: Chinese

AIM: To intensively investigate the chemical constituents of cultured (3y) Taxus chinensis var. mairei. METHODS: The whole plant was extracted with ethanol, the ethanol extract was subjected to extraction with methylene chloride, which was submitted to chromatog. on silica gel and Sephadex column to isolate some compds. And their structures were elucidated on the basis of spectral anal. (UV, IR, ESI-MS, 1H NMR, 13C NMR). RESULTS: The compds. were identified as 2α -deacetoxytaxinine J, taxuyunnanine C, yunnanxane, taxinine J, 1-dehydroxy-baccatin VI, taxol, 19-debenzoyl-19-acetyltaxinine M, taxinine M, taxicin, taxa-4(20)-11-diene- 2α , 5α , 10β -triacetoxy- 14β , 2-methybulyrate, baccatin III, cephalomannine, 7,13-dideacetyl-9,10-didebenzoyltaxchinin C7,13-dideacetyl-9,10-didebenzoyltaxchinin C, taxamairin A, taxamairin B, α -conidendrin, secoisolariciresinol, isotaxiresinol, $\beta\text{-sitosterol}$ and daucosterol; 19-debenzoyl-19-acetyltaxinine M and C7,13-dideacetyl-9,10-didebenzoyltaxchinin C were isolated from this source for the first time. CONCLUSION: The chemical constituents of planted Taxus mairei are nearly the same as that of the wild.

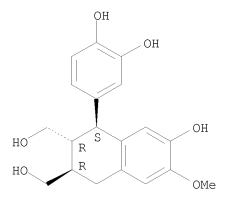
RL: NPO (Natural product occurrence); PRP (Properties); BIOL (Biological study); OCCU (Occurrence)

(chemical constituents of cultured Taxus mairei)

26194-57-0 CAPLUS RN

2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-CN hydroxy-6-methoxy-, (1S, 2R, 3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



L13 ANSWER 16 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

2008:1226505 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 150:370166

TITLE: Chemical constituents in heartwood of Taxus

vunnanensis

AUTHOR(S): Chen, Xueying; Liang, Jingyu

CORPORATE SOURCE: Department of Phytochemistry, China Pharmaceutical

University, Nanjing, Jiangsu Province, 210009, Peop.

Rep. China

SOURCE: Zhongcaoyao (2007), 38(7), 979-982

CODEN: CTYAD8; ISSN: 0253-2670

PUBLISHER: Zhongcaoyao Zazhi Bianjibu

DOCUMENT TYPE: Journal LANGUAGE: Chinese

The chemical constituents in the heartwood of Taxus yunnanensis

Cheng et L. K. Fu were intensively investigated. The heartwood was extracted with ethanol, and ethanol extract was subjected to the extraction with

methylene

trichloride, which was submitted to chromatog. on silica gel and Sephadex LH-20 column to isolate some compds. And their structures were elucidated on the basis of spectral anal. (UV, IR, ESI-MS, 1H-NMR and 13C-NMR). The compds. were identified as

 2α , 5α , 7β , 9α , 10β , 13α -hexaacetoxy-4(20), 11-

taxadiene (I), taxusin (II), taxa-4(20),11-diene-

 2α , 5α , 10β -triacetoxy- 14β , 2-methybutyrate (III),

 10β -hydroxy- 2α , 5α , 14β -triacetoxy-4(20), 11-taxadiene

(IV), 1-dehydroxybaccatin IV (V), baccatin IV (VI), baccatin VI (VII),

7,9-deacetylbaccatin VI (VIII), 10-deacetyltaxuyannine (IX),

 $1\beta\text{-acetoxy-}5\text{-deacetyl-baccatin I (X), baccatin I (XI), taxuchin A}$

(XII), secoisolariciresinol (XIII), α -conidendrin (XIV), isotaxiresinol (XV), lariciresinol (XVI), sequoyitol (XVII) and

 $\beta\text{-sitosterol}$ (XVIII). Among them compds. I, V, VI, XI, XII and XIV were obtained from the heartwood of T. yunnanensis for the first time. conclusion, the chemical constituents in the heartwood differed from the

other parts of T. yunnanensis, but there was little difference within the species of Taxus L.

IT 26194-57-0P, Isotaxiresinol

RL: BSU (Biological study, unclassified); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); PREP (Preparation) (chemical constituents in heartwood of **Taxus** yunnanensis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 17 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1999:693513 CAPLUS

DOCUMENT NUMBER: 132:33212

TITLE: Lignans, flavonoids and phenolic derivatives from

Taxus mairei

AUTHOR(S): Yang, Shung-Jim; Fang, Jim-Min; Cheng, Yu-Shia

CORPORATE SOURCE: Department of Chemistry, National Taiwan University,

Taipei, 106, Taiwan

SOURCE: Journal of the Chinese Chemical Society (Taipei)

(1999), 46(5), 811-818

CODEN: JCCTAC; ISSN: 0009-4536

PUBLISHER: Chinese Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

AB From the twigs of **Taxus** mairei, 35 lignans, 2 sesquilignans, 4 flavonoids, 3 bisflavonoids, 13 phenolic derivs., 2 sesquiterpenes, 3 bisnorsesquiterpenes, 3 long-chain carboxylic acids and 4 steroids were isolated. The new lignans and phenolic glucosides include

7'-hydroxynortrachelogenin, 7-hydroxymatairesinol,

3'-O-demethylepipinoresinol, taxiresinol 9-acetate, 3'-O-demethyltanegool,

8'-epitanegool, 3,3'-dimethoxy-4,4',9-trihydroxy-7,9'-epoxylignan-7'-one,

3-O-demethyldihydrodehydrodiconiferyl alc., taxumaiglucoside A

heptaacetate, taxumaiglucoside B heptaacetate, and taxumaiglucoside C heptaacetate. Their structures were determined by spectral methods.

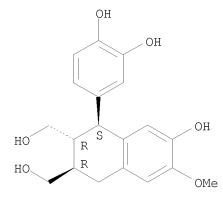
IT **26194-57-0**, (+)-Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(from **Taxus** mairei)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)



REFERENCE COUNT: 70 THERE ARE 70 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 18 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2007:66436 CAPLUS

DOCUMENT NUMBER: 146:291589

TITLE: Antiallergic activity of aqueous extracts and

constituents of Taxus yunnanensis

AUTHOR(S): Koyama, Junko; Morita, Izumi; Kobayashi, Norihiro;

Hirai, Keiichi; Simamura, Eriko; Nobukawa, Takahiro;

Kadota, Shiqetoshi

CORPORATE SOURCE: Kobe Pharmaceutical University, Higashinada-ku, Kobe,

658-8558, Japan

SOURCE: Biological & Pharmaceutical Bulletin (2006), 29(11),

2310-2312

CODEN: BPBLEO; ISSN: 0918-6158

PUBLISHER: Pharmaceutical Society of Japan

DOCUMENT TYPE: Journal LANGUAGE: English

GΙ

The H2O, H2O/MeOH (1: 1) exts. from the wood of **Taxus** yunnanensis showed a remarkable inhibitory effect on induced histamine release from the human basophilic cell line, KU812. The eleven constituents purified from the wood exts. of **Taxus** yunnanensis were tested by an in vitro histamine release inhibition assay. Among them, secoisolarciresinol and taxiresinol were found to show inhibitory activities. A new neolignan, 2-[2-hydroxy-5-(3-hydroxypropyl)-3-methoxyphenyl]-1-(4-hydroxy-3-methoxyphenyl)propane-1,3-diol (I), was isolated from the wood of **Taxus** yunnanensis.

IT 26194-57-0P, Isotaxiresinol

RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(antiallergic activity of aqueous exts. and constituents of ${\bf Taxus}$ yunnanensis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 19 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:334697 CAPLUS

DOCUMENT NUMBER: 146:77986

TITLE: Hypoglycemic effects of the wood of Taxus

yunnanensis on streptozotocin-induced diabetic rats

and its active components

AUTHOR(S): Banskota, A. H.; Nguyen, N. T.; Tezuka, Y.; Nobukawa,

T.; Kadota, S.

CORPORATE SOURCE: Institute of Natural Medicine, Toyama Medical and

Pharmaceutical University, 2630-Sugitani, Toyama,

930-0194, Japan

SOURCE: Phytomedicine (2006), 13(1-2), 109-114

CODEN: PYTOEY; ISSN: 0944-7113

PUBLISHER: Elsevier GmbH

DOCUMENT TYPE: Journal LANGUAGE: English

AB Hypoglycemic effects of the H2O and MeOH exts. of the wood of

Taxus yunnanensis were examined in streptozotocin (STZ)-induced
diabetic rats. The H2O extract significantly lowered the fasting blood
glucose level by 33.7% at a 100 mg/kg dose on i.p. administration. From
the active H2O extract of the wood, three lignans, i.e., isotaxiresinol (1),
secoisolariciresinol (2) and taxiresinol (3), were isolated as major
components. These lignans were further tested for their hypoglycemic
effects on the same exptl. model. At a dose of 100 mg/kg (i.p.),
isotaxiresinol (1) reduced the fasting blood glucose level of diabetic
rats by 34.5%, while secoisolariciresinol (2) and taxiresinol (3) reduced
by 33.4% and 20.9%, resp. The blood glucose lowering effects of 1 and 2
were stronger than the mixture of tolbutamide (200 mg/kg) and buformin (1
mg/kg) used as a pos. control, which lowered fasting blood glucose level
by 24.0%.

IT 26194-57-0P, Isotaxiresinol

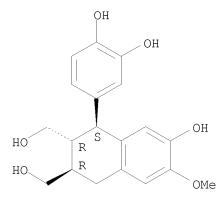
RL: BSU (Biological study, unclassified); NPO (Natural product occurrence); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses) (TLC, HPLC of water extract from **Taxus** yunnanensis wood show

isotaxiresinol, secoisolariciresinol, taxiresinol have varied hypoglycemic effect on streptozotocin-induced diabetic rat)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 20 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2003:555111 CAPLUS

DOCUMENT NUMBER: 140:70921

TITLE: DPPH radical scavenging and nitric oxide inhibitory

activities of the constituents from the wood of

Taxus yunnanensis

AUTHOR(S): Banskota, Arjun H.; Tezuka, Yasuhiro; Nguyen, Nhan

Trung; Awale, Suresh; Nobukawa, Takahiro; Kadota,

Shigetoshi

CORPORATE SOURCE: Institute of Natural Medicine, Toyama Medical and

Pharmaceutical University, Toyama, Japan

SOURCE: Planta Medica (2003), 69(6), 500-505 CODEN: PLMEAA; ISSN: 0032-0943

PUBLISHER: Georg Thieme Verlag

DOCUMENT TYPE: Journal

LANGUAGE: English AB The H2O, H2O/MeOH (1:1) and MeOH exts. of the wood of ${\bf Taxus}$

yunnanensis possessed significant DPPH radical scavenging and nitric oxide (NO) inhibitory activities. Chemical investigation of these exts. led us to

isolation of nineteen compds., i.e., five lignans, two simple phenolics, and twelve taxane-type diterpenes. Isotaxiresinol and seco-

isolariciresinol, two major lignans of the wood, possessed potent

DPPH radical scavenging activities with IC50 values of 21.7 and 28.9

 $\mu\text{M}\text{,}$ resp. Similarly, coniferyl aldehyde, taxusin,

10-deacetyltaxuyunnanine C, hongdoushan A, and

 2α , 5α , 10β -triacetoxy- 14β -[(S)-2-methylbutyryloxy]-

4(20),11- taxadiene showed potent NO inhibitory activity with IC50 values of 18.0, 22.1, 28.5, 15.0 and 26.4 $\mu\text{M},$ resp., which were either equal or lower than the pos. control NG-monomethyl-L-arginine (L-NMMA) with an

IC50 value of 28.5 $\mu\text{M}.$ IT 26194-57-0P, Isotaxiresinol

RL: NPO (Natural product occurrence); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

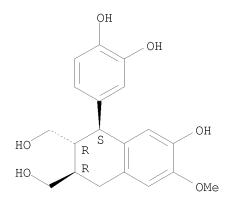
(DPPH scavenging and NO inhibitory activities of constituents from

Taxus yunnanensis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 21 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1994:692369 CAPLUS

DOCUMENT NUMBER: 121:292369

ORIGINAL REFERENCE NO.: 121:53195a,53198a

TITLE: Hypoglycemic and antiplatelet constituents of

Taxus mairei

AUTHOR(S): Guo, Daih-Huang; Ko, Huey-Ming; Lai, Jem-min; Chiu,

Tai-Hui; Wu, Tian-Shung; Teng, Che-Ming; Kuo,

Sheng-Chu

CORPORATE SOURCE: Department Pharmacy, Tajen Junior College Pharmacy

Pingtong, Taiwan

SOURCE: Chinese Pharmaceutical Journal (Taipei, Taiwan)

(1994), 46(3), 175-83

CODEN: CPHJEP; ISSN: 1016-1015

DOCUMENT TYPE: Journal LANGUAGE: English

AB Bioassay-directed fractionation of the BuOH extract of the heartwood of T. mairei led to the isolation of isotaxiresinol (I). I showed significant hypoglycemic effect at 100 mg/Kg in rat and with weak antiplatelet aggregation effect. In addition, (-)secoisolariciresinol , vanillin and $\beta\text{-sitostenone}$ from the chloroform extract also revealed antiplatelet aggregation activity.

IT **26194-57-0**, Isotaxiresinol

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(hypoglycemic and blood platelet aggregation inhibitory constituents of **Taxus** mairei)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

L13 ANSWER 22 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1990:175629 CAPLUS

DOCUMENT NUMBER: 112:175629

ORIGINAL REFERENCE NO.: 112:29611a,29614a

TITLE: Reinvestigation on the constituents of the heartwood

of Taiwan yew

AUTHOR(S): Chuang, Li Chin; Chen, Kwei Ju; Lin, Yun Shan; Chen,

Fa Ching

CORPORATE SOURCE: Dep. Chem., Tamkang Univ., Tamsui, Taiwan

SOURCE: Taiwan Kexue (1989), 42, 29-35 CODEN: TKHSAU; ISSN: 0015-7791

DOCUMENT TYPE: Journal LANGUAGE: Chinese

AB 1-Dehydroxybaccatin IV and (+)-dihydroquercetin (taxifolin) were isolated

from Taxus mairei in addition to a long-chain alkane, long-chain

ester, β -sitosterol, taxusin,

taxa-4(20), 11-diene- 5α , 9α , 10β , 13α -tetrol-

 9α , 10β -diacetate, taxa-4(20), 11-diene- 2α , 5α , 7β , 10β -tetrol- 5α , 7β , 10β -

triacetate- 2α - α -methylbutyrate, secoisolariciresinol, and

isotaxiresinol.

IT **26194-57-0**, Isotaxiresinol

RL: BIOL (Biological study)

(of Taiwan yew heartwood)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

L13 ANSWER 23 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1997:770200 CAPLUS

DOCUMENT NUMBER: 128:86475

ORIGINAL REFERENCE NO.: 128:16825a, 16828a

TITLE: A lignan from roots of **Taxus** mairei

AUTHOR(S): Shen, Ya-Ching; Chen, Ching-Yeu; Lin, Yat-Min; Kuo,

Yao-Haur

CORPORATE SOURCE: Institute of Marine Resources. National. Sun Yat-sen

University, Kaohsiung, Taiwan

SOURCE: Phytochemistry (1997), 46(6), 1111-1113

CODEN: PYTCAS; ISSN: 0031-9422

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

GΙ

AB A new lignan, taxumairin (I) was isolated from the roots of Formosan **Taxus** mairei, along with known lignans. The structure of taxumairin has been characterized as (+)-7,8-trans-8,8'-trans-7',8'-trans-7-(3-methoxy-4-hydroxy)phenyl-7'-(3'-methoxy-4'-hydroxy)phenyl-8-hydroxymethyl-8'-ethoxymethyltetrahydrofuran, on the basis of spectral analyses.

IT **26194-57-0**, Isotaxiresinol

RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
BIOL (Biological study); OCCU (Occurrence)
 (from roots of Taxus mairei)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 24 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1998:389753 CAPLUS

DOCUMENT NUMBER: 129:180010

ORIGINAL REFERENCE NO.: 129:36477a,36480a

TITLE: Bioactive lignans and taxoids from the roots of

Formosan Taxus mairei

AUTHOR(S): Shen, Ya-Ching; Chen, Ching-Yeu; Chen, Yin-Ju; Kuo,

Yao-Haur; Chien, Ching-Te; Lin, Yat-Min

CORPORATE SOURCE: Institute Marine Resources, National Sun Yat-sen

University, Kaohsiung, Taiwan

SOURCE: Chinese Pharmaceutical Journal (Taipei) (1997),

49 (5-6), 285-296

CODEN: CPHJEP; ISSN: 1016-1015

PUBLISHER: Pharmaceutical Society of Republic of China

DOCUMENT TYPE: Journal LANGUAGE: English

AB Four lignans, (-)- α -conidendrin, (-)-secoisolariciresinol, isotaxiresinol, and taxiresinol and 4 taxoids, 1β -dehydroxybaccatin VI, 1β -dehydroxybaccatin IV, 1β -hydroxybaccatin I and taxumairol D were isolated from the roots of Formosan T. mairei. The structures of

these compds. were established by spectral and chemical anal. The lignans exhibited potent cytotoxicities against KB-16, A-549 and HT-29 tumor cells.

cells.

IT **26194-57-0**, Isotaxiresinol

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(bioactive lignans and taxoids from Taxus mairei riits)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 25 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2003:867253 CAPLUS

DOCUMENT NUMBER: 140:209916

TITLE: Absolute configuration and anticancer activity of

taxiresinol and related lignans of Taxus

wallichiana

AUTHOR(S): Chattopadhyay, Sunil K.; Kumar, T. R. Santha; Maulik,

Prakas R.; Srivastava, Sachin; Garg, Ankur; Sharon, Ashoke; Negi, Arvind S.; Khanuja, Suman Preet S.

CORPORATE SOURCE: Central Institute of Medicinal and Aromatic Plants

(CIMAP), PO CIMAP, Lucknow, 226 015, India

SOURCE: Bioorganic & Medicinal Chemistry (2003), 11(23),

4945-4948

CODEN: BMECEP; ISSN: 0968-0896

PUBLISHER: Elsevier Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Absolute configuration of taxiresinol 1, a lignan from the heartwood of **Taxus** wallichiana has been determined as 8R, 8'R, and 7'R with the help

of chemical correlation method and x-ray crystallog. The anticancer activity of taxiresinol 1 and other two lignans 2, 3 were also studied.

Taxiresinol 1 showed notable anticancer activity in the in vitro bioassays

against colon, liver, ovarian and breast cancer cell lines.

IT 26194-57-OP, Isotaxiresinol

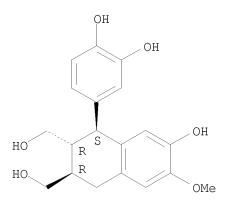
RL: PAC (Pharmacological activity); PRP (Properties); PUR (Purification or recovery); RCT (Reactant); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

(absolute configuration and anticancer activity of taxiresinol and related lignans of **Taxus** wallichiana)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 26 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:823549 CAPLUS

DOCUMENT NUMBER: 143:186763

TITLE: Therapeutic/preventive agent for osteoporosis

containing as component isotaxiresinol derived from

Taxus yunnanensis

INVENTOR(S): Kadota, Shigetoshi; Nobukawa, Takahiro

PATENT ASSIGNEE(S): Kotosugi Inc., Japan SOURCE: PCT Int. Appl., 17 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| P | ATENT | NO. | KIND DATE | | | | | APPL | ICAT | DATE | | | | | | | | | |
|------------------------|---------------|------|-----------|-----|------|------|------------|------|------|------------------|------|------|-----|-----|----------|------|-----|--|--|
| W(| WO 2005074905 | | | | | | 20050818 | | | WO 2005-JP1055 | | | | | 20050127 | | | | |
| | W: | ΑE, | AG, | AL, | AM, | ΑT, | ΑU, | ΑZ, | BA, | BB, | BG, | BR, | BW, | BY, | BZ, | CA, | CH, | | |
| | | CN, | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC, | EE, | EG, | ES, | FI, | GB, | GD, | | |
| | | GE, | GH, | GM, | HR, | HU, | ID, | IL, | IN, | IS, | JP, | ΚE, | KG, | KP, | KR, | KΖ, | LC, | | |
| | | LK, | LR, | LS, | LT, | LU, | LV, | MA, | MD, | MG, | MK, | MN, | MW, | MX, | MZ, | NA, | NΙ, | | |
| | | NO, | NΖ, | OM, | PG, | PH, | PL, | PT, | RO, | RU, | SC, | SD, | SE, | SG, | SK, | SL, | SY, | | |
| | | ТJ, | TM, | TN, | TR, | TT, | TZ, | UA, | UG, | US, | UZ, | VC, | VN, | YU, | ZA, | ZM, | ZW | | |
| | RW: | BW, | GH, | GM, | ΚE, | LS, | MW, | ΜZ, | NA, | SD, | SL, | SZ, | TZ, | UG, | ZM, | ZW, | ΑM, | | |
| | | ΑZ, | BY, | KG, | KΖ, | MD, | RU, | ТJ, | TM, | ΑT, | BE, | BG, | CH, | CY, | CZ, | DE, | DK, | | |
| | | EE, | ES, | FI, | FR, | GB, | GR, | HU, | ΙE, | IS, | ΙΤ, | LT, | LU, | MC, | NL, | PL, | PT, | | |
| | | RO, | SE, | SI, | SK, | TR, | BF, | ВJ, | CF, | CG, | CI, | CM, | GΑ, | GN, | GQ, | GW, | ML, | | |
| | | MR, | NE, | SN, | TD, | TG | | | | | | | | | | | | | |
| CI | CN 1842327 | | | | | | 2006 | 1004 | | CN 2005-80000941 | | | | | 20050127 | | | | |
| $T\Gamma$ | TW 285108 | | | | | | 2007 | 0811 | | TW 2005-94103196 | | | | | 20050202 | | | | |
| KI | KR 2006037416 | | | | | | A 20060503 | | | KR 2006-702230 | | | | | 20060201 | | | | |
| PRIORITY APPLN. INFO.: | | | | | | | | | | JP 2004-26535 | | | | | A 2 | 0040 | 203 | | |
| | | | | | | | | | | WO 2 | 005- | JP10 | 55 | 1 | W 2 | 0050 | 127 | | |
| OTHER S | SOURCE | (S): | MAR: | PAT | 143: | 1867 | 63 | | | | | | | | | | | | |

OTHER SOURCE(S):

AΒ A medicine useful in treatments for and prevention of osteoporosis. The medicine for treatments for and prevention of osteoporosis contains as an active ingredient a compound represented by the formula (I; R1 represents C1-4 alkyloxy) or a medically acceptable salt or ester of the compound of the formula I. Of the compds. represented by I, the compound wherein R1 is CH30 is isotaxiresinol derived from Taxus yunnanensis. This compound functions to inhibit bone absorption and has physiol. activity in accelerating bone formation.

26194-57-ODP, Isotaxiresinol, derivs. and salts ΙT 26194-57-0P, Isotaxiresinol

RL: PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(Therapeutic/preventive agent for osteoporosis containing as component isotaxiresinol derived from Taxus yunnanensis)

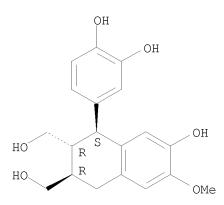
RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 27 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:80489 CAPLUS

DOCUMENT NUMBER: 140:117348

TITLE: Hypoglycemic agent, liver protecting agent and anticancer agent containing lignans originating in

Taxus yunnanensis

INVENTOR(S): Kadota, Shigetoshi; Nobukawa, Takahiro

PATENT ASSIGNEE(S): Kotosugi Inc., Japan SOURCE: PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT N | KIN | D | DATE | | | APPL | DATE | | | | | | | | | |
|---------------|-----|-----|------|-------------|-----|------|------|------|------|-----|----------|-----|-----|-----|-----|-----|
| WO 2004009065 | | | | A1 20040129 | | | ; | WO 2 | 003- | | 20030723 | | | | | |
| W: | ΑE, | AG, | AL, | AM, | ΑT, | ΑU, | ΑZ, | BA, | BB, | BG, | BR, | BY, | BZ, | CA, | CH, | CN, |
| | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC, | EE, | ES, | FΙ, | GB, | GD, | GE, | GH, |
| | GM, | HR, | HU, | ID, | IL, | IN, | IS, | JP, | ΚE, | KG, | KP, | KR, | KΖ, | LC, | LK, | LR, |
| | LS, | LT, | LU, | LV, | MA, | MD, | MG, | MK, | MN, | MW, | MX, | MZ, | NI, | NO, | NZ, | OM, |

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PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     AU 2003248097
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                                 20040209
                                            AU 2003-248097
                                                                    20030723
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                                            US 2005-522186
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                                20080222
                                            HK 2005-109274
                                                                    20051020
                          Α1
     KR 2006086458
                                20060731
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                          Α
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                                                                    20080208
PRIORITY APPLN. INFO.:
                                             JP 2002-214694
                                                                 A 20020724
                                             JP 2003-119178
                                                                 A 20030424
                                             WO 2003-JP9370
                                                                 W 20030723
                                             KR 2005-701127
                                                                 A3 20050121
                                             US 2005-522186
                                                                 A3 20050124
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OTHER SOURCE(S): MARPAT 140:117348

AB Disclosed are drugs containing taxiresinol, (7'R)-7' -hydroxylariciresinol, secoisolariciresinol and isotaxiresinol, which are lignans contained in hongdoushan (**Taxus** yunnanensis), as the active ingredients.

Drugs contains an extract, which is obtained by extracting a hongdoushan plant with water and further extracting the obtained extract with an organic solvent, as

the active ingredient. These drugs are useful particularly as a hyopglycermic agent, a liver protecting agent and an anticancer agent.

IT **26194-57-0P**, Isotaxiresinol

RL: PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(hypoglycemic agent, liver protecting agent and anticancer agent containing lignans originating in **Taxus** yunnanensis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 28 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:334692 CAPLUS

DOCUMENT NUMBER: 145:241621

TITLE: In vivo anti-osteoporotic activity of isotaxiresinol,

a lignan from wood of **Taxus** yunnanensis

AUTHOR(S): Yin, J.; Tezuka, Y.; Subehan; Shi, L.; Nobukawa, M.;

Nobukawa, T.; Kadota, S.

CORPORATE SOURCE: Institute of Natural Medicine, Toyama Medical and

Pharmaceutical University, 2630 Sugitani, Toyama,

930-0194, Japan

SOURCE: Phytomedicine (2006), 13(1-2), 37-42

CODEN: PYTOEY; ISSN: 0944-7113

PUBLISHER: Elsevier GmbH

DOCUMENT TYPE: Journal LANGUAGE: English

AB Isotaxiresinol, the main lignan isolated from the water extract of wood of **Taxus** yunnanensis, was investigated for its effect on bone loss, on serum biochem. markers for bone remodeling and on uterine tissue, using ovariectomized (OVX) rats as the model of postmenopausal osteoporosis. After oral administration of isotaxiresinol (50 and 100 mg/kg/d) for 6 wk, bone mineral content (BMC) and bone mineral d. (BMD) in total and cortical bones were increased as compared to those of OVX control rats, and decreases of three bone strength indexes induced by OVX surgery were prevented. Serum biochem. markers for bone remodeling revealed that isotaxiresinol slightly increased bone formation and significantly inhibited bone resorption without side effect on uterine tissue. These results suggest that isotaxiresinol may be useful for treatment of postmenopausal osteoporosis, especially for prevention of bone fracture induced by estrogen deficiency.

IT 26194-57-0P, Isotaxiresinol

RL: NPO (Natural product occurrence); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(lignan from T. yunnanensis isotaxiresinol showed anti-osteoporotic activity by increasing BMC and BMD in cortical bone with higher bone formation and inhibited bone resorption in uterine tissue of rat model of postmenopausal osteoporosis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 29 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1975:533681 CAPLUS

DOCUMENT NUMBER: 83:133681

ORIGINAL REFERENCE NO.: 83:21038h,21039a

TITLE: Inhibitory effect of **Taxus** mairei heartwood

extractives on the curing of unsaturated polyester

resins

AUTHOR(S): Lee, Chuen Lai; Hirose, Yoshiyuki; Nakatsuka,

Tomoichiro

CORPORATE SOURCE: Fac. Agric., Univ. Tokyo, Tokyo, Japan Mokuzai Gakkaishi (1975), 21(4), 249-56

CODEN: MKZGA7; ISSN: 0021-4795

DOCUMENT TYPE: Journal LANGUAGE: Japanese

AB Among 5 compds. extracted from **Taxus** mairei heartwood, i.e., sequoyitol (I) [523-92-2], taxusin (II) [19605-80-2], α -conidendrin

(III) [518-55-8], secoisolariciresinol (IV) [25327-50-8], and isotaxiresinol (V) [**26194-57-0**], only V inhibited curing of

polyester resin in the presence of Bz202, and IV and V inhibited the

curing in the presence of MeCOEt peroxide and Co salts.

IT 26194-57-0P

RL: PREP (Preparation)

(from **Taxus** mairei, crosslinking inhibitors for unsatd.

polyester)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

L13 ANSWER 30 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2009:44496 CAPLUS

DOCUMENT NUMBER: 150:275688

TITLE: Development of an LC-ESI-MS/MS method for the

determination of histamine: Application to the

quantitative measurement of histamine degranulation by

KU812 cells

AUTHOR(S): Koyama, Junko; Takeuchi, Atsuko; Tode, Chisato;

Shimizu, Maki; Morita, Izumi; Nobukawa, Machiko;

Nobukawa, Makiko; Kobayashi, Norihiro

CORPORATE SOURCE: Kobe Pharmaceutical University, Higashinada, Kobe,

658-8558, Japan

SOURCE: Journal of Chromatography, B: Analytical Technologies

in the Biomedical and Life Sciences (2009), 877(3),

207-212

CODEN: JCBAAI; ISSN: 1570-0232

PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

A rapid, simple, and sensitive liquid chromatog.-electrospray ionization AΒ tandem mass spectrometry (LC-ESI-MS/MS) method was developed for the identification and quantification of histamine without a previous derivatization step or the addition of general ion-pairing reagents to the mobile phase. This method was used to measure histamine release following degranulation of KU812 human basophilic cells, using pyrazol as an internal standard Analyses were performed on an LC system employing a Cosmosil 5C18 PAQ column and an isocratic elution with methanol-0.005% trifluoroacetic acid (1:1) at a flow rate of 0.2 mL/min. A triple-quadrupole mass spectrometer, equipped with an electrospray ionization interface was employed, operating in the pos. ion mode. retention time of histamine and the internal standard were 4.0 and 5.0 min, resp. The relative standard deviations (R.S.D.s) of the retention time and peak area were between 0.47% and 2.03%. Micropipette tip solid-phase extraction (SPE) using LooseTip C18 allowed for not only rapid sample preparation,

but also decreased suppression effects, improving peak shape. This method was used to evaluate the anti-allergic effects of compds. contained in **Taxus** yunnanensis exts. Four constituents that were isolated from the wood exts. of T. yunnanensis and sodium cromoglicate, which is used as a first line anti-allergic drug, were tested in an in vitro histamine release inhibition assay. Of these compds., taxiresinol and isotaxiresinol were more inhibitory than sodium cromoglicate.

IT **26194-57-0**, Isotaxiresinol

RL: PAC (Pharmacological activity); BIOL (Biological study) (development of LC-ESI-MS/MS method for determination of histamine following degranulation of KU812 human basophilic cells and use in evaluating anti-allergic effects of compds. contained in **Taxus** yunnanensis exts.)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 31 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:243556 CAPLUS

DOCUMENT NUMBER: 140:350498

TITLE: Secoisolariciresinol and isotaxiresinol inhibit tumor

necrosis factor- α -dependent hepatic apoptosis in

mice

AUTHOR(S): Banskota, Arjun H.; Nguyen, Nhan Trung; Tezuka,

Yasuhiro; Le Tran, Quan; Nobukawa, Takahiro;

Kurashige, Youichi; Sasahara, Masakiyo; Kadota,

Shigetoshi

CORPORATE SOURCE: Institute of Natural Medicine, Toyama Medical and

Pharmaceutical University, Toyama, 930-0194, Japan

SOURCE: Life Sciences (2004), 74(22), 2781-2792

CODEN: LIFSAK; ISSN: 0024-3205

PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English

The effects of secoisolariciresinol (1) and isotaxiresinol (2), two major lignans isolated from the wood of Taxus yunnanensis, on tumor necrosis factor- α (TNF- α)-dependent hepatic apoptosis induced by D-galactosamine (D-GalN)/lipopolysaccharide (LPS) were investigated in mice. Co-administration of d-GalN (700 mg/kg) and LPS (10 μ g/kg) resulted in a typical hepatic apoptosis characterized by DNA fragmentation and the formation of apoptotic bodies. Serum glutamic pyruvic transaminase (sGPT) and glutamic oxaloacetic transaminase (sGOT) levels were also raised at 8 h after D-GalN/LPS intoxication due to a severe necrosis of hepatocytes. Pre-administration of 1 or 2 (50, 10 mg/kg, i.p.) 12 and 1 h before d-GalN/LPS significantly reduced DNA fragmentation and prevented chromatin condensation, apoptotic body formation and hepatitis. Pro-inflammatory cytokines such as TNF- α and interferon- γ (IFN- γ) secreted from LPS-activated macrophages are important mediators of hepatocyte apoptosis in this model. Pre-treatment with 1 or 2 significantly inhibited the elevation of serum $TNF-\alpha$ and $IFN-\gamma$ levels. In a sep. experiment, both lignans had a significant dose-dependent protective effect on D-GalN/TNF-lpha-induced cell death in primary cultured mouse hepatocytes and $TNF-\alpha$ -mediated cell death in murine L929 fibrosarcoma cells. These results indicated that 1 and 2 prevent D-GalN/LPS-induced hepatic injury by inhibiting hepatocyte apoptosis through the blocking of TNF- α and IFN- γ production by activated macrophages and direct inhibition of the apoptosis induced by TNF- α .

IT **26194-57-0**, Isotaxiresinol

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(secoisolariciresinol and isotaxiresinol inhibit tumor necrosis factor- α -dependent hepatic apoptosis in mice)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

35

L13 ANSWER 32 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN

2001:417499 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 135:166155

TITLE: In Vitro Metabolism of Plant Lignans: New Precursors

of Mammalian Lignans Enterolactone and Enterodiol AUTHOR(S): Heinonen, Satu; Nurmi, Tarja; Liukkonen, Kirsi; Poutanen, Kaisa; Waehaelae, Kristiina; Deyama,

Takeshi; Nishibe, Sansei; Adlercreutz, Herman

CORPORATE SOURCE: Folkhaelsan Research Center and Department of Clinical

Chemistry, University of Helsinki, Helsinki,

FIN-00014, Finland

SOURCE: Journal of Agricultural and Food Chemistry (2001),

49(7), 3178-3186

CODEN: JAFCAU; ISSN: 0021-8561

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

The metabolism of the plant lignans matairesinol, secoisolariciresinol, pinoresinol, syringaresinol, arctigenin, 7-hydroxymatairesinol, isolariciresinol, and lariciresinol by human fecal microflora was investigated to study their properties as mammalian lignan precursors. The quant. analyses of lignan precursors and the mammalian lignans enterolactone and enterodiol were performed by HPLC with coulometric electrode array detector. The metabolic products, including mammalian lignans, were characterized as trimethylsilyl derivs. by gas chromatog.-mass spectrometry. Matairesinol, secoisolariciresinol, lariciresinol, and pinoresinol were converted to mammalian lignans only. Several metabolites were isolated and tentatively identified as for syringaresinol and arctigenin in addition to the mammalian lignans. Metabolites of 7-hydroxymatairesinol were characterized as enterolactone and 7-hydroxyenterolactone by comparison with authentic reference compds. A

metabolic scheme describing the conversion of the most abundant new mammalian lignan precursors, pinoresinol and lariciresinol, is presented.

ΙT 477-72-5

AB

RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL (Biological study); FORM (Formation, nonpreparative)

(plant lignans as precursors of mammalian lignans enterolactone and enterodiol)

RN 477-72-5 CAPLUS

2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-CN hydroxy-6-methoxy- (CA INDEX NAME)

THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 32 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L13 ANSWER 11 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN
     2007:1012507 CAPLUS
    147:465158
DN
    Antioxidant activity of polyphenols from the far-east plant Taxus
TΙ
     cuspidata
    Veselova, M. V.; Fedoreev, S. A.; Vasilevskaya, N. A.; Denisenko, V. A.;
ΑU
     Gerasimenko, A. V.
     Pacific Institute of Bioorganic Chemistry, Far-East Division, Russian
CS
     Academy of Sciences, Vladivostok, Russia
     Pharmaceutical Chemistry Journal (2007), 41(2), 88-93
SO
    CODEN: PCJOAU; ISSN: 0091-150X
PВ
     Springer
DT
    Journal
    English
LA
RE.CNT 24
              THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L13 ANSWER 1 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN
ΑN
     1952:54579 CAPLUS
DN
     46:54579
OREF 46:9086b-i,9087a-b
     Isotaxiresinol (3'-dimethylisolariciresinol), a new lignan extracted from
TT
     the heartwood of the English yew, Taxus
     baccata
     King, F. E.; Jurd, L.; King, T. J.
ΑU
CS
     Univ. Nottingham, UK
SO
     Journal of the Chemical Society (1952) 17-24
    CODEN: JCSOA9; ISSN: 0368-1769
    Journal
DT
    Unavailable
LA
    CASREACT 46:54579
OS
L13 ANSWER 2 OF 32 CAPLUS COPYRIGHT 2009 ACS on STN
ΑN
    1973:111031 CAPLUS
    78:111031
DN
OREF 78:17819a,17822a
     Taxiresinol, a new lignan in the heartwood of Taxus
     baccata
    Mujumdar, R. B.; Srinivasan, R.; Venkataraman, K.
ΑU
CS
    Natl. Chem. Lab., Poona, India
SO
    Indian Journal of Chemistry (1972), 10(7), 677-80
    CODEN: IJOCAP; ISSN: 0019-5103
DΤ
    Journal
    English
LA
=> d l11 1-2 ibib abs hitstr
L11 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN
                         2006:334692 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         145:241621
TITLE:
                         In vivo anti-osteoporotic activity of
                         isotaxiresinol, a lignan from wood of Taxus
                         yunnanensis
AUTHOR(S):
                         Yin, J.; Tezuka, Y.; Subehan; Shi, L.; Nobukawa, M.;
                         Nobukawa, T.; Kadota, S.
CORPORATE SOURCE:
                         Institute of Natural Medicine, Toyama Medical and
```

Pharmaceutical University, 2630 Sugitani, Toyama,

930-0194, Japan

SOURCE: Phytomedicine (2006), 13(1-2), 37-42

CODEN: PYTOEY; ISSN: 0944-7113

PUBLISHER: Elsevier GmbH

DOCUMENT TYPE: Journal LANGUAGE: English

AB Isotaxiresinol, the main lignan isolated from the water extract of wood of

Taxus yunnanensis, was investigated for its effect on **bone** loss, on serum biochem. markers for **bone** remodeling and on uterine

tissue, using ovariectomized (OVX) rats as the model of postmenopausal

osteoporosis. After oral administration of isotaxiresinol (50 and

100 mg/kg/d) for 6 wk, **bone** mineral content (BMC) and **bone** mineral d. (BMD) in total and cortical **bones** were

increased as compared to those of OVX control rats, and decreases of three

bone strength indexes induced by OVX surgery were prevented. Serum biochem. markers for **bone** remodeling revealed that

isotaxiresinol slightly increased **bone** formation and significantly inhibited **bone resorption** without side

effect on uterine tissue. These results suggest that isotaxiresinol may be useful for treatment of postmenopausal **osteoporosis**, especially for prevention of **bone** fracture induced by estrogen deficiency.

IT 26194-57-0P, Isotaxiresinol

RL: NPO (Natural product occurrence); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(lignan from T. yunnanensis isotaxiresinol showed anti-

osteoporotic activity by increasing BMC and BMD in cortical

 $\begin{tabular}{ll} \textbf{bone} & with higher & \textbf{bone} & formation and inhibited \\ \end{tabular}$

bone resorption in uterine tissue of rat model of

postmenopausal osteoporosis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:823549 CAPLUS

DOCUMENT NUMBER: 143:186763

TITLE: Therapeutic/preventive agent for **osteoporosis**

containing as component isotaxiresinol derived from

Taxus yunnanensis

INVENTOR(S): Kadota, Shigetoshi; Nobukawa, Takahiro

PATENT ASSIGNEE(S): Kotosugi Inc., Japan

SOURCE: PCT Int. Appl., 17 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PA. | TENT : | KIND | | DATE | | APPLICATION NO. | | | | | | DATE | | | | | | |
|------------------------|---------------|------|------|------|-----|-----------------|------|----------|-----|------------------|------------------|------|-----|-----|-----|----------|-----|--|
| WO | WO 2005074905 | | | | | A1 | | 20050818 | | WO 2005-JP1055 | | | | | | | | |
| | W: | ΑE, | AG, | AL, | ΑM, | ΑT, | ΑU, | ΑZ, | BA, | BB | , BG, | BR, | BW, | BY, | ΒZ, | CA, | CH, | |
| | | CN, | CO, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ | , EC, | EE, | EG, | ES, | FΙ, | GB, | GD, | |
| | | GE, | GH, | GM, | HR, | HU, | ID, | IL, | IN, | IS, | , JP, | ΚE, | KG, | KP, | KR, | KΖ, | LC, | |
| | | LK, | LR, | LS, | LT, | LU, | LV, | MA, | MD, | MG. | , MK, | MN, | MW, | MX, | MZ, | NA, | NI, | |
| | | NO, | NΖ, | OM, | PG, | PH, | PL, | PT, | RO, | RU | , SC, | SD, | SE, | SG, | SK, | SL, | SY, | |
| | | ΤJ, | TM, | TN, | TR, | TT, | TZ, | UA, | UG, | US, | , UZ, | VC, | VN, | YU, | ZA, | ZM, | ZW | |
| | RW: | BW, | GH, | GM, | KE, | LS, | MW, | MΖ, | NA, | SD | , SL, | SZ, | TZ, | UG, | ZM, | ZW, | ΑM, | |
| | | ΑZ, | BY, | KG, | KΖ, | MD, | RU, | ТJ, | TM, | AT, | , BE, | BG, | CH, | CY, | CZ, | DE, | DK, | |
| | | EE, | ES, | FΙ, | FR, | GB, | GR, | ΗU, | ΙE, | IS, | , IT, | LT, | LU, | MC, | NL, | PL, | PT, | |
| | | RO, | SE, | SI, | SK, | TR, | BF, | ВJ, | CF, | CG, | , CI, | CM, | GΑ, | GN, | GQ, | GW, | ML, | |
| | | MR, | ΝE, | SN, | TD, | ΤG | | | | | | | | | | | | |
| CN | CN 1842327 | | | | | | 2006 | 1004 | | CN 2005-80000941 | | | | | | 20050127 | | |
| TW | TW 285108 | | | | | | 2007 | 20070811 | | | IW 2005-94103196 | | | | 2 | 0050 | 202 | |
| KR | KR 2006037416 | | | | | | 2006 | 0503 | | KR 2 | 2006- | 7022 | 30 | | 2 | 0060 | 201 | |
| PRIORITY APPLN. INFO.: | | | | | | | | | | JP 2 | 2004- | 2653 | 5 | | A 2 | 0040 | 203 | |
| | | | | | | | | | | WO 2 | 2005- | JP10 | 55 | | W 2 | 0050 | 127 | |
| OTHER SO | MAR: | PAT | 143: | 1867 | 63 | | | | | | | | | | | | | |

GΙ

AB A medicine useful in treatments for and prevention of osteoporosis

. The medicine for treatments for and prevention of osteoporosis
contains as an active ingredient a compound represented by the formula (I;
R1 represents C1-4 alkyloxy) or a medically acceptable salt or ester of
the compound of the formula I. Of the compds. represented by I, the compound
wherein R1 is CH3O is isotaxiresinol derived from Taxus yunnanensis. This
compound functions to inhibit bone absorption and has physiol.
activity in accelerating bone formation.

IT 26194-57-ODP, Isotaxiresinol, derivs. and salts 26194-57-OP, Isotaxiresinol

Ι

RL: PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(Therapeutic/preventive agent for **osteoporosis** containing as component isotaxiresinol derived from Taxus yunnanensis)

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

RN 26194-57-0 CAPLUS

CN 2,3-Naphthalenedimethanol, 1-(3,4-dihydroxyphenyl)-1,2,3,4-tetrahydro-7-hydroxy-6-methoxy-, (1S,2R,3R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).

8

REFERENCE COUNT:

THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT